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IN THE CLAIMS:

Please amend the claims as follows:

- (Amended) A humanized antibody that binds ICAM-1, said antibody selected from: SEQ ID NO:1 and 3 (HumA); SEQ ID NO:5 and 7 (HumB); SEQ ID NO:9 and 11 (HumC); SEQ ID NO:13 and 15 (HumD); SEQ ID NO:17 and 19 (HumE); SEQ ID NO:21 and 23 (HumF); SEQ ID NO:25 and 27 (HumG); SEQ ID NO:29 and 31 (HumH); and SEQ ID NO:33 and 35 (HumI).
- 2. (Amended) A subsequence of the antibody of claim 1, said antibody subsequence capable of binding an ICAM-1 epitope, and wherein the variable framework region of said subsequence has one or more amino acids of a human consensus variable framework region sequence.
- 3. (Amended) The humanized antibody subsequence of claim 2, wherein the antibody subsequence comprises a single chain, Fab, Fab' or (Fab)₂ fragment.
- 4. (Amended) The humanized antibody of claim 1, said antibody having one or more amino acid substitutions, provided that said <u>substitution</u> when in the variable framework region is an <u>amino acid based upon a human consensus variable framework region sequence, and said substituted antibody is capable of binding an ICAM-1 epitope.</u>
- 5. (Amended) A humanized antibody that binds ICAM-1 and inhibits pathogen infection of cells expressing ICAM-1, wherein the variable framework region has one or more amino acids of a human consensus variable framework region sequence, and wherein the protective efficacy is at least equivalent to mouse monoclonal antibody denoted as 1A6.
- 6. (Previously Presented) The humanized antibody of claim 5, said antibody having a protective efficacy at least 2 times greater than the non-humanized antibody.
- 7. (Previously Presented) The humanized antibody of claim 5, said antibody having a protective efficacy at least 5 times greater than the non-humanized antibody.
- 8. (Previously Presented) The humanized antibody of claim 5, said antibody having a protective efficacy at least 10 times greater than the non-humanized antibody.
- 9. (Previously Presented) The humanized antibody of claim 5, said antibody having a protective efficacy at least 20 times greater than the non-humanized antibody.
- 10. (Previously Presented) The humanized antibody of claim 5, said antibody having a protective efficacy at least 30 times greater than the non-humanized antibody.

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11. (Previously Presented) The humanized antibody of claim 5, wherein the pathogen is human rhinovirus (HRV).

- 12. (Amended) The humanized antibody of claim 5, wherein the pathogen is eoxackie coxsackie A virus, or respiratory syncytial virus, or malaria.
- 13. (Previously Presented) The humanized antibody of claim 5, wherein the antibody is an intact immunoglobulin molecule comprising 2 full-length heavy chains and 2 full-length light chains.
- 14. (Previously Presented) The humanized antibody of claim 5, wherein the antibody is an antibody subsequence that binds to ICAM-1.
- 15. (Previously Presented) The humanized antibody of claim 14, wherein the antibody subsequence comprises a single chain, Fab, Fab' or (Fab)₂ fragment.
- 16. (Amended) The humanized antibody of claim 1 or 5, wherein the antibody is multispecific or multifunctional.[(;)]
- 17. (Amended) The humanized antibody of claim 1 or 5, wherein the antibody is linked to one or more identical or different antibodies to form a multimer.
- 18. (Previously Presented) The humanized antibody of claim 17, wherein the multimer comprises a homo- or hetero-dimer, trimer, or tetramer.
- 19. (Previously Presented) The humanized antibody of claim 17, wherein the multimer is formed via a multimerization domain.
- 20. (Previously Presented) The humanized antibody of claim 19, wherein the multimerization domain comprises a human amino acid sequence.
- 21. (Previously Presented) The humanized antibody of claim 19, further comprising a linker located between the multimerization domain and the antibody.
- 22. (Amended) A humanized antibody that inhibits human rhinovirus (HRV) infection of cells comprising the amino acid sequence set forth in any of SEQ ID NO:1 and 3 (HumA); SEQ ID NO:5 and 7 (HumB); SEQ ID NO:9 and 11 (HumC); SEQ ID NO:13 and 15 (HumD); SEQ ID NO:17 and 19 (HumE); SEQ ID NO:21 and 23 (HumF); SEQ ID NO:25 and 27 (HumG); SEQ ID NO:29 and 31 (HumH); and SEQ ID NO:33 and 35 (HumI); or a subsequence thereof, wherein the variable framework region of said subsequence has one or more amino acids of a human consensus variable framework region sequence.

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23. (Previously Presented) The humanized antibody of claim 22, wherein the antibody is an immunoglobulin molecule comprising 2 full-length heavy chain polypeptides and 2 full-length light chain polypeptides.

- 24. (Previously Presented) The humanized antibody of claim 22, wherein the subsequence comprises a single chain, Fab, Fab' or (Fab)₂ fragment.
- 25. (Previously Presented) The humanized antibody of claim 22, wherein the antibody is linked with other identical or different antibodies to form a multimer.
- 26. (Previously Presented) The humanized antibody of claim 25, wherein the multimer comprises a homo- or hetero-dimer, trimer, or tetramer.
- 27. (Previously Presented) The humanized antibody of claim 25, wherein the different antibodies are human, humanized or non-human.
- 28. (Amended) A nucleic acid sequence encoding a humanized antibody of claim 1 or 22 or a subsequence thereof, wherein the variable framework region of said subsequence has one or more amino acids of a human consensus variable framework region sequence.
- 29. (Previously Presented) An expression cassette comprising the nucleic acid sequence of claim 28 operably linked to an expression control element.
- 30. (Previously Presented) A vector comprising the nucleic acid sequence of claim 29.
- 31. (Previously Presented) The vector of claim 29, wherein the nucleic acid sequence is operably linked to an expression control element.
- 32. (Previously Presented) A cell comprising the nucleic acid sequence of claim 28.
- 33. (Previously Presented) The cell of claim 31, wherein the cell is prokaryotic or eukaryotic.
- 34. (Previously Presented) A pharmaceutical composition comprising a humanized antibody of claim 1 or 5, and a pharmaceutically acceptable carrier.
- 35. (Previously Presented) The pharmaceutical composition of claim 34, wherein the carrier is compatible with inhalation or nasal delivery to a subject.
- 36. (Previously Presented) A method of inhibiting pathogen infection of a cell comprising contacting a pathogen or a cell with an amount of a humanized antibody of claims 1 or 5, sufficient to inhibit pathogen infection of the cell.
- 37. (Previously Presented) The method of claim 36, wherein the cell is present in a subject.
- 38. (Previously Presented) The method of claim 37, wherein the cell is an epithelial cell.
- 39. (Previously Presented) The method of claim 37, wherein the cell expresses ICAM-1.

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40. (Amended) A method of inhibiting HRV infection of a cell comprising contacting HRV or a cell susceptible to HRV infection with an amount of a humanized antibody of claim [21] 22 effective to inhibit HRV infection of the cell.

- 41. (Previously Presented) The method of claim 40, wherein the cell is present in a subject.
- 42. (Previously Presented) The method of claim 41, wherein the subject has or is at risk of having asthma.
- 43. (Previously Presented) The method of claim 40, wherein the antibody binds to an antigen present on the surface of the cell.
- 44. (Previously Presented) The method of claim 40, wherein the cell expresses ICAM-1.
- 45. (Previously Presented) The method of claim 40, wherein the cell is an epithelial cell.
- 46. (Previously Presented) The method of claim 40, wherein the humanized antibody is administered locally.
- 47. (Amended) The method of claim 40, wherein the humanized antibody is administered via inhalation or intranasaly intranassaly.
- 48. (Amended) A method of inhibiting HRV infection, inhibiting HRV progression or treating HRV infection of a subject comprising administering to a subject having or at risk of having HRV infection an amount of a humanized antibody of claim 21 22 effective to inhibit, inhibit progression or treat HRV infection of the subject.
- 49. (Previously Presented) The method of claim 48, wherein the humanized antibody is administered locally.
- 50. (Amended) The method of claim 48, wherein the humanized antibody is administered via inhalation or intranasaly intranassaly.
- 51. (Previously Presented) The method of claim 48, wherein the subject has or is at risk of having asthma.
- 52. (Previously Presented) The method of claim 48, wherein the subject is a newborn or between the ages of 1 to 5, 5 to 10 or 10 to 18.
- 53. (Amended) A method of decreasing or inhibiting one or more symptoms of the common cold in a subject comprising administering to a subject having a common cold an amount of a humanized antibody of claim 21 22 effective to decrease or inhibit one or more symptoms of the common cold in the subject.

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54. (Previously Presented) The method of claim 53, wherein the humanized antibody is administered locally.

- 55. (Amended) The method of claim 53, wherein the humanized antibody is administered via inhalation or intranasaly intranassaly.
- 56. (Previously Presented) The method of claim 53, wherein the subject has or is at risk of having asthma.
- 57. (Previously Presented) The method of claim 53, wherein the subject is a newborn or between the ages of 1 to 5, 5 to 10 or 10 to 18.

Please add the following new claims:

- 58. (New) The humanized antibody of claim 4, wherein said substitution is in a complementarity determining region.
- 59. (New) The humanized antibody of claim 4, wherein said substitution is in a framework region.
- 60. (New) The humanized antibody of claim 4, wherein said substitution comprises 5-10 amino acids.
- 61. (New) The humanized antibody of claim 4, wherein said substitution comprises 3-5 amino acids.
- 62. (New) The humanized antibody of claim 4, wherein said substitution comprises 1-3 amino acids.
- 63. (New) The humanized antibody of claim 4, wherein the antibody binds ICAM-1 with increased affinity relative to humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 64. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 4-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 65. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 5-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.

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66. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 5 to 8-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.

- 67. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 5 to 10-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 68. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 8 to 15-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 69. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 10 to 20-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 70. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 20 to 40-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 71. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 50 to 100-fold greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 72. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 100-fold or greater than humanized antibody having the complementarity regions of mouse monoclonal antibody denoted as 1A6.
- 73. (New) The humanized antibody of claim 4, wherein the antibody binds ICAM-1 with an affinity at least equivalent to mouse monoclonal antibody denoted as 1A6.
- 74. (New) The humanized antibody of claim 4, wherein the antibody binds ICAM-1 with increased affinity relative to mouse monoclonal antibody denoted as 1A6.
- 75. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 4-fold greater than mouse monoclonal antibody denoted as 1A6.
- 76. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 5- fold greater than mouse monoclonal antibody denoted as 1A6.
- 77. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 5 to 8-fold greater than mouse monoclonal antibody denoted as 1A6.

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78. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 5 to 10-fold greater than mouse monoclonal antibody denoted as 1A6.

- 79. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 8 to 15-fold greater than mouse monoclonal antibody denoted as 1A6.
- 80. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 10 to 20-fold greater than mouse monoclonal antibody denoted as 1A6.
- 81. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 20 to 40-fold greater than mouse monoclonal antibody denoted as 1A6.
- 82. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 50 to 100-fold greater than mouse monoclonal antibody denoted as 1A6.
- 83. (New) The humanized antibody of claim 4, wherein the antibody has an ICAM-1 binding affinity 100-fold or greater than mouse monoclonal antibody denoted as 1A6.